Preliminary Synthesis of Climate Change Impacts on O_3'' is available primarily via the Internet on the National Center for Environmental Assessment's home page under the Recent Additions and the Data and Publications menus at *www.epa.gov/ncea.* A limited number of paper copies are available from the Information Management Team, NCEA; *telephone:* 703–347–8561; *facsimile:* 703–347–8691. If you are requesting a paper copy, please provide your name, mailing address, and the document title. Copies are not available from Versar, Inc.

FOR FURTHER INFORMATION CONTACT: Questions regarding information, registration, access or services for individuals with disabilities, or logistics for the external peer-review workshop should be directed to Versar, Inc., 6850 Versar Center, Springfield, VA 22151; *telephone:* 703–750–3000, extension 737 or 211; *facsimile: Attn.:* Malikah Moore at 703–642–6954; e-mail: *mmoore@versar.com* and *dsinkowski@versar.com.* To request accommodation of a disability, please contact Malikah Moore at *mmoore@versar.com* or 703–750–3000, ext. 211, preferably at least 10 days prior to the meeting, to give as much time as possible to process your request.

If you need technical information about the document, please contact Brooke L. Hemming, Ph.D., National Center for Environmental Assessment (NCEA); *telephone:* 919–541–5668; *facsimile:* 919–541–7885; *e-mail: hemming.brooke@epa.gov.*

SUPPLEMENTARY INFORMATION:

I. Summary of Information About the Project/Document

The ''Interim Report of the U.S. EPA Global Change Research Program Assessment of the Impacts of Global Change on Regional U.S. Air Quality: A Preliminary Synthesis of Climate Change Impacts on O_3'' is intended to provide air quality managers and scientists a summary and synthesis of the scientific results that have emerged from the EPA ORD Global Change Research Program (ORD GCRP) assessment of the impact of global change on U.S. regional air quality. The report discusses the studies that have focused on the sensitivity of U.S. air quality to meteorological changes associated with a warming climate in large regions within the continental U.S. The EPA recognizes that climate-air quality interactions occur at multiple scales (both spatial and temporal), and that an understanding of these interactions demands contributions from several scientific disciplines. The EPA ORD GCRP developed a research and assessment program that combines the resources and expertise of the ORD labs and centers toward the goal of developing the necessary scientific underpinnings. The ultimate goal of the Program is to provide air quality managers with the scientific information and tools needed to evaluate the implications of global change and to enhance their ability to consider global change in their decisions. The ''Interim Report of the U.S. EPA Global Change Research Program Assessment of the Impacts of Global Change on Regional U.S. Air Quality: A Preliminary Synthesis of Climate Change Impacts on O_3'' is a preliminary step in that direction. This report provides an update of the progress that has been made in applying climate and atmospheric chemistry models to investigate potential future meteorological effects on air quality. It does not include changes in air pollutant emissions other than those that are explicitly linked to meteorological variables and incorporated within the models (*e.g.*, biogenic VOC emissions). In addition, it provides a preliminary interpretation of what this improved scientific understanding means for air quality management. Future assessment reports will cover the combined impacts of changing climate and air pollutant emissions on air quality. The program also plans to develop additional reports that focus on additional pollutants, including PM and mercury. NCEA worked collaboratively with the EPA Office of Air and Radiation (OAR), and ORD's National Risk Management Research Laboratory (NRMRL), National Exposure Research Laboratory (NERL) and National Center for Environmental Research (NCER) on this report.

II. Workshop Information

Members of the public may attend the workshop as observers, and there will be a limited time for comments from the public in the afternoon. Please let Versar, Inc. know if you wish to make comments during the workshop. Space is limited, and reservations will be accepted on a first-come, first-served basis.

Dated: July 25, 2008.

Rebecca Clark,

Deputy Director, National Center for Environmental Assessment.

[FR Doc. E8–17925 Filed 8–4–08; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2008–0545; FRL–8375–9]

Pesticides; Revised Fee Schedule for Registration Applications

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: EPA is publishing a revised list of pesticide registration service fees applicable to specified pesticide applications and tolerance actions. Under the Pesticide Registration Improvement Renewal Act, the registration service fees for covered pesticide registration applications received on or after October 1, 2008, increase by 5% rounding up to the nearest dollar from the fees published for fiscal year 2008, and certain decision review periods have been reduced. The new fees and decision review periods become effective on October 1, 2008.

FOR FURTHER INFORMATION CONTACT: Elizabeth Leovey (7501P), Immediate Office, Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460–0001; telephone number: (703) 305–7328; fax number: (703) 308–4776; e-mail address: *leovey.elizabeth@epa.gov.*

SUPPLEMENTARY INFORMATION:

I. General Information

A. Does this Action Apply to Me?

You may be potentially affected by this action if you register pesticide products under the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA). Potentially affected entities may include, but are not limited to:

- Agricultural pesticide manufacturers (32532).
- Antimicrobial pesticide manufacturers (32561).
- Antifoulant pesticide manufacturers (32551).
- Wood preservative manufacturers (32519).

This listing is not intended to be exhaustive, but rather provides a guide for readers regarding entities likely to be affected by this action. Other types of entities not listed in this unit could also be affected. The North American Industrial Classification System (NAICS) codes have been provided to assist you and others in determining whether this action might apply to certain entities. To determine whether you or your business may be affected by this action, you should carefully examine the applicability provisions in the notice and in FIFRA section 33. If

you have any questions regarding the applicability of this action to a particular entity, consult the person listed under **FOR FURTHER INFORMATION CONTACT**.

B. How Can I Get Copies of this Document and Other Related Information?

1. *Docket*. EPA has established a docket for this action under docket identification (ID) number EPA–HQ–OPP–2008–[0545]. Publicly available docket materials are available either in the electronic docket at *http://www.regulations.gov*, or, if only available in hard copy, at the Office of Pesticide Programs (OPP) Regulatory Public Docket in Rm. S–4400, One Potomac Yard (South Bldg.), 2777 S. Crystal Dr., Arlington, VA. The hours of operation of this Docket Facility are from 8:30 a.m. to 4 p.m., Monday through Friday, excluding legal holidays. The Docket Facility telephone number is (703) 305–5805.

2. *Electronic access*. You may access this **Federal Register** document electronically through the EPA Internet under the ''**Federal Register**'' listings at *http://www.epa.gov/fedrgstr*.

II. Background

A. What Action is the Agency Taking?

The Pesticide Registration Improvement Act of 2003 established a new section 33 of FIFRA creating a registration service fee system for certain types of pesticide applications, establishment of tolerances, and certain other regulatory decisions under FIFRA and the Federal Food, Drug, and Cosmetic Act (FFDCA). Section 33 also created a schedule of decision review times for applications covered by the service fee system. The Agency began administering the registration service fee system for covered applications received on or after March 23, 2004.

On October 9, 2007, the Pesticide Registration Improvement Renewal Act was signed by the President, revising, among other things, FIFRA section 33. The new law reauthorized the service fee system through 2012 and established fees and review times for applications received during fiscal years 2008 through 2012. As required by section 33(b)(6)(A) of FIFRA, the registration service fees for covered pesticide registration applications received on or after October 1, 2008, increase by five percent rounding up to the nearest dollar from the fees published in the October 30, 2007, **Federal Register** Notice (72 FR 61466). In addition, this notice provides shorter decision review periods for certain application categories.

B. What is the Agency's Authority for Taking this Action?

The publication of this fee schedule is required by section 33(b)(6)(C) of FIFRA as amended.

III. Elements of the Fee Schedule

This unit explains how EPA has organized the fee schedule identified in the statute and how to read the fee schedule tables, and includes a key to terminology published with the table in the Congressional Review. EPA's organization and presentation of the fee schedule information does not affect the categories of registration service fees or the structure or procedures for submitting applications or petitions for tolerance.

A. The Congressional Record Fee Schedule

The fee schedule published in the Congressional Record of July 21, 2007 identifies the registration service fees and decision times and is organized according to the organizational units of the Office of Pesticide Programs (OPP) within EPA. Thereafter, the categories within the organizational unit sections of the table are further categorized according to the type of application being submitted, the use patterns involved, or, in some cases, upon the type of pesticide that is the subject of the application. The fee categories differ by Division. Not all application types are covered by, or subject to, the fee system.

B. Fee Schedule and Decision Review Times

In today's notice, EPA has retained the format of previous schedule notices and included the corrections to the schedule published in the September 24, 2007 issue of the Congressional Record. The schedules are presented as 11 tables, organized by OPP Division and by type of application or pesticide subject to the fee. These tables only list the decision time review periods for fiscal years 2009 and 2010 as these are the only applicable review periods for applications received on or after October 1, 2008. Unit IV. presents fee tables for the Registration Division (RD) (5 tables), the Antimicrobials Division (AD) (3 tables), and the Biopesticides and Pollution Prevention Division (BPPD) (3 tables).

C. How to Read the Tables

1. *Each table consists of the following columns*:

- The column entitled ''EPA No.'' assigns an EPA identifier to each fee category. There are 140 categories spread across the 3 Divisions. There are 58 RD categories, 27 AD categories, and 55 BPPD categories. For tracking purposes, OPP has assigned a 3-digit identifier to each category, beginning with RD categories, followed by AD and BPPD categories. The categories are prefaced with a letter designation indicating which Division of OPP is responsible for applications in that category (R= Registration Division, A=Antimicrobials Division, B=Biopesticides and Pollution Prevention Division).
- The column entitled ''CR No.'' cross-references the current Congressional Record category number for convenience. However, EPA will be using the categories as numbered in the ''EPA No.'' column in its tracking systems.
- The column entitled ''Action'' describes the categories of action. In establishing the expanded fee schedule categories, Congress eliminated some of the more confusing terminology of the original categories. For example, instead of the term ''fast-track,'' the schedule in the Congressional Record uses the regulatory phrase ''identical or substantially similar in composition and use to a registered product.''
- The column entitled ''Decision Time'' lists the decision times in months for each type of action for Fiscal Years 2009 and 2010. The 2010 decision times apply to 2011 and 2012. The decision review periods in the tables are based upon EPA fiscal years (FY), which run from October 1 through September 30.
- The column entitled ''FY 09/FY 10 Registration Service Fee ($)'' lists the registration service fee for the action for fiscal year 2009 (October 1, 2008 through September 30, 2009) and fiscal year 2010 (October 1, 2009 through September 30, 2010).

2. *The following acronyms are used in some of the tables*:

- DART–Dose Adequacy Response Team
- DNT–Developmental Neurotoxicity
- HSRB–Human Studies Review Board
- GW/SW–Ground Water/Surface Water
- PHI–Pre-Harvest Interval
- PPE–Personal Protective Equipment
- REI–Restricted Entry Interval
- SAP–FIFRA Scientific Advisory Panel

IV. PRIRA Fee Schedule Tables—Effective October 1, 2008

A. Registration Division (RD)

The Registration Division of OPP is responsible for the processing of pesticide applications and associated tolerance petitions for pesticides that are termed ''conventional chemicals,'' excluding pesticides intended for antimicrobial uses. The term ''conventional chemical'' is a term of art that is intended to distinguish synthetic chemicals from those that are of naturally occurring or non-synthetic origin, synthetic chemicals that are identical to naturally-occurring chemicals and microbial pesticides. Tables 1 through 5 of Unit IV.A. cover RD actions.

TABLE 1.—REGISTRATION DIVISION—NEW ACTIVE INGREDIENTS

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
R010	1	Food use[1]	24	24	542,115
R020	2	Food use; reduced risk[1]	18	18	542,115
R030	3	Food use; Experimental Use Permit application submitted simultaneously with application for registration; decision time for Experimental Use Permit and temporary tolerance same as #R040[1]	24	24	599,235
R040	4	Food use; Experimental Use Permit application; establish temporary tolerance; submitted before application for registration; credit $326,025 toward new active ingredient application that follows	18	18	399,525
R050	5	Food use; application submitted after Experimental Use Permit application; decision time begins after Experimental Use Permit and temporary tolerance are granted[1]	14	14	199,815
R060	6	Non-food use; outdoor[1]	21	21	376,635
R070	7	Non-food use; outdoor; reduced risk[1]	16	16	376,635
R080	8	Non-food use; outdoor; Experimental Use Permit application submitted simultaneously with application for registration; decision time for Experimental Use Permit same as #R090[1]	21	21	416,640
R090	9	Non-food use; outdoor; Experimental Use Permit application submitted before application for registration; credit $228,225 toward new active ingredient application that follows	16	16	279,615
R100	10	Non-food use; outdoor; submitted after Experimental Use Permit application; decision time begins after Experimental Use Permit is granted[1]	12	12	137,025
R110	11	Non-food use; indoor[1]	20	20	209,475
R120	12	Non-food use; indoor; reduced risk[1]	14	14	209,475
R121	13	Non-food use; indoor; Experimental Use Permit application submitted before application for registration; credit $100,000 toward new active ingredient application that follows	18	18	157,500
R122	14	Enriched isomer(s) of registered mixed-isomer active ingredient[1]	18	18	273,945
R123	15	Seed treatment only; includes non-food and food uses; limited uptake into Raw Agricultural Commodities[1]	18	18	407,610
R124	16	Conditional Ruling on Preapplication Study Waivers; applicant-initiated	6	6	2,184

[1]All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.

TABLE 2.—REGISTRATION DIVISION—NEW USES

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
R130	17	First food use; indoor; food/food handling[1]	21	21	165,375
R140	18	Additional food use; indoor; food/food handling	15	15	38,588
R150	19	First food use[1]	21	21	228,270
R160	20	First food use; reduced risk[1]	16	16	228,270
R170	21	Additional food use	15	15	57,120
R180	22	Additional food use; reduced risk	10	10	57,120
R190	23	Additional food uses; 6 or more submitted in one application	15	15	342,720
R200	24	Additional food uses; 6 or more submitted in one application; reduced risk	10	10	342,720
R210	25	Additional food use; Experimental Use Permit application; establish temporary tolerance; no credit toward new use registration	12	12	42,315
R220	26	Additional food use; Experimental Use Permit application; crop destruct basis; no credit toward new use registration	6	6	17,136
R230	27	Additional use; non-food; outdoor	15	15	22,827
R240	28	Additional use; non-food; outdoor; reduced risk	10	10	22,827
R250	29	Additional use; non-food; outdoor; Experimental Use Permit application; no credit toward new use registration	6	6	17,136
R260	30	New use; non-food; indoor	12	12	11,025
R270	31	New use; non-food; indoor; reduced risk	9	9	11,025
R271	32	New use; non-food; indoor; Experimental Use Permit application; no credit toward new use registration	6	6	8,400
R272	33	Review of Study Protocol; applicant-initiated; excludes DART, pre-registration conferences, Rapid Response review, DNT protocol review, protocols needing HSRB review	3	3	2,184
R273	34	Additional use; seed treatment; limited uptake into Raw Agricultural Commodities; includes crops with established tolerances (e.g., for soil or foliar application); includes food or non-food uses	12	12	43,575
R274	35	Additional uses; seed treatment only; 6 or more submitted in one application; limited uptake into Raw Agricultural Commodities; includes crops with established tolerances (e.g., for soil or foliar application); includes food and/or non-food uses	12	12	261,450

[1]All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.

TABLE 3.—REGISTRATION DIVISION—IMPORT AND OTHER TOLERANCES

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
R280	36	Establish import tolerance; new active ingredient or first food use[1]	21	21	275,625
R290	37	Establish import tolerance; additional food use	15	15	55,125
R291	38	Establish import tolerances; additional food uses; 6 or more crops submitted in one petition	15	15	330,750
R292	39	Amend an established tolerance (e.g., decrease or increase); domestic or import; applicant-initiated	10	10	39,165
R293	40	Establish tolerance(s) for inadvertent residues in one crop; applicant-initiated	12	12	46,200
R294	41	Establish tolerances for inadvertent residues; 6 or more crops submitted in one application; applicant-initiated	12	12	277,200
R295	42	Establish tolerance(s) for residues in one rotational crop in response to a specific rotational crop application; applicant-initiated	15	15	57,120
R296	43	Establish tolerances for residues in rotational crops in response to a specific rotational crop petition; 6 or more crops submitted in one application; applicant-initiated	15	15	342,720

[1]All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.

TABLE 4.—REGISTRATION DIVISION—NEW PRODUCTS

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
R300	44	New product; identical or substantially similar in composition and use to a registered product; no data review or only product chemistry data; cite-all data citation, or selective data citation where applicant owns all required data, or applicant submits specific authorization letter from data owner. Category also includes 100% re-package of registered end-use or manufacturing-use product that requires no data submission nor data matrix.	3	3	1,365
R301	45	New product; identical or substantially similar in composition and use to a registered product; registered source of active ingredient; selective data citation only for data on product chemistry and/or acute toxicity and/or public health pest efficacy, where applicant does not own all required data and does not have a specific authorization letter from data owner.	4	4	1,638
R310	46	New end-use or manufacturing-use product; requires review of data package within RD; includes reviews and/or waivers of data for only: • Product chemistry and/or • Acute toxicity and/or • Public health pest efficacy	6	6	4,578
R311	49	New product; requires approval of new food-use inert; applicant-initiated; excludes approval of safeners	12	12	16,317
R312	50	New product; requires approval of new non-food-use inert; applicant-initiated	6	6	8,715

TABLE 4.—REGISTRATION DIVISION—NEW PRODUCTS—Continued

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
R313	51	New product; requires amendment to existing inert tolerance exemption (e.g., adding post-harvest use); applicant-initiated	10	10	11,991
R320	47	New product; new physical form; requires data review in science divisions	12	12	11,424
R330	48	New manufacturing-use product; registered active ingredient; selective data citation	12	12	17,136
R331	52	New product; repack of identical registered end-use product as a manufacturing-use product; same registered uses only	3	3	2,184
R332	53	New manufacturing-use product; registered active ingredient; unregistered source of active ingredient; submission of completely new generic data package; registered uses only	24	24	244,650

TABLE 5.—REGISTRATION DIVISION—AMENDMENTS TO REGISTRATION

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
R340	54	Amendment requiring data review within RD (e.g., changes to precautionary label statements, or source changes to an unregistered source of active ingredient)[1]	4	4	3,444
R350	55	Amendment requiring data review in science divisions (e.g., changes to REI, or PPE, or PHI, or use rate, or number of applications; or add aerial application; or modify GW/SW advisory statement)[1]	8	8	11,424
R370	56	Cancer reassessment; applicant-initiated	18	18	171,255
R371	57	Amendment to Experimental Use Permit; requires data review/risk assessment	6	6	8,715
R372	58	Refined ecological and/or endangered species assessment; applicant-initiated	18	12	163,065

[1]EPA-initiated amendments shall not be charged fees. Fast-track amendments handled by the Antimicrobials Division are to be completed within the FIFRA stated timelines listed in section 3(h) and are not subject to PRIA fees. Label amendments submitted by notification under PR Notices, such as PR Notice 95–2 and PR Notice 98–10, continue under PR Notice timelines and are not subject to PRIA fees.

B. Antimicrobials Division (AD)

The Antimicrobials Division of OPP is responsible for the processing of pesticide applications and associated tolerances for conventional chemicals intended for antimicrobial uses, that is, uses that are defined under FIFRA section 2(mm)(1)(A), including products for use against bacteria, protozoa, non-agricultural fungi, and viruses. AD is also responsible for a selected set of conventional chemicals intended for other uses, including most wood preservatives and antifoulants. Tables 6 through 8 of Unit IV.B. cover AD actions.

TABLE 6.—ANTIMICROBIALS DIVISION—NEW ACTIVE INGREDIENTS

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
A380	59	Food use; establish tolerance exemption[1]	24	24	99,225
A390	60	Food use; establish tolerance[1]	24	24	165,375
A400	61	Non-food use; outdoor; FIFRA section 2(mm) uses[1]	18	18	82,688

TABLE 6.—ANTIMICROBIALS DIVISION—NEW ACTIVE INGREDIENTS—Continued

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
A410	62	Non-food use; outdoor; uses other than FIFRA section 2(mm)[1]	21	21	165,375
A420	63	Non-food use; indoor; FIFRA section 2(mm) uses[1]	18	18	55,125
A430	64	Non-food use; indoor; uses other than FIFRA section 2(mm)[1]	20	20	82,688
A431	65	Non-food use; indoor; low-risk and low-toxicity food-grade active ingredient(s); efficacy testing for public health claims required under GLP and following DIS/TSS or AD-approved study protocol	12	12	57,750

[1]All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.

TABLE 7.—ANTIMICROBIALS DIVISION—NEW USES

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
A440	66	First food use; establish tolerance exemption[1]	21	21	27,563
A450	67	First food use; establish tolerance[1]	21	21	82,688
A460	68	Additional food use; establish tolerance exemption	15	15	11,025
A470	69	Additional food use; establish tolerance	15	15	27,563
A480	70	Additional use; non-food; outdoor; FIFRA section 2(mm) uses	9	9	16,538
A490	71	Additional use; non-food; outdoor; uses other than FIFRA section 2(mm)	15	15	27,563
A500	72	Additional use; non-food; indoor; FIFRA section 2(mm) uses	9	9	11,025
A510	73	Additional use; non-food; indoor; uses other than FIFRA section 2(mm)	12	12	11,025
A520	74	Experimental Use Permit application	9	9	5,513
A521	75	Review of public health efficacy study protocol within AD; per AD Internal Guidance for the Efficacy Protocol Review Process; applicant-initiated; Tier 1	4	3	2,100
A522	76	Review of public health efficacy study protocol outside AD by members of AD Efficacy Protocol Review Expert Panel; applicant-initiated; Tier 2	15	12	10,500

[1]All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.

TABLE 8.—ANTIMICROBIALS DIVISION—NEW PRODUCTS AND AMENDMENTS

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
A530	77	New product; identical or substantially similar in composition and use to a registered product; no data review or only product chemistry data; cite-all data citation, or selective data citation where applicant owns all required data, or applicant submits specific authorization letter from data owner. Category also includes 100% re-package of registered end-use or manufacturing-use product that requires no data submission nor data matrix.	3	3	1,103
A531	78	New product; identical or substantially similar in composition and use to a registered product; registered source of active ingredient; selective data citation only for data on product chemistry and/or acute toxicity and/or public health pest efficacy, where applicant does not own all required data and does not have a specific authorization letter from data owner	4	4	1,575
A532	85	New product; identical or substantially similar in composition and use to a registered product; registered active ingredient; unregistered source of active ingredient; cite-all data citation except for product chemistry; product chemistry data submitted	4	4	4,410
A540	79	New end-use product; FIFRA section 2(mm) uses only	4	4	4,410
A550	80	New end-use product; uses other than FIFRA section 2(mm); non-FQPA product	6	6	4,410
A560	81	New manufacturing-use product; registered active ingredient; selective data citation	12	12	16,538
A570	82	Label amendment requiring data submission[1]	4	4	3,308
A571	83	Cancer reassessment; applicant-initiated	18	18	82,688
A572	84	Refined ecological risk and/or endangered species assessment; applicant-initiated	18	12	78,750

[1]EPA-initiated amendments shall not be charged fees. Fast-track amendments handled by the Antimicrobials Division are to be completed within the FIFRA stated timelines listed in section 3(h) and are not subject to PRIA fees. Label amendments submitted by notification under PR Notices, such as PR Notice 95–2 and PR Notice 98–10, continue under PR Notice timelines and are not subject to PRIA fees.

C. Biopesticides and Pollution Prevention Division (BPPD)

The Biopesticides and Pollution Prevention Division of OPP is responsible for the processing of pesticide applications for biochemical pesticides, microbial pesticides, and plant-incorporated protectants (PIPs).

The fee tables for BPPD tables are presented by type of pesticide rather than by type of action: Microbial and biochemical pesticides, straight chain lepidopteran pheromones (SCLPs), and PIPs. Within each table, the types of application are the same as those in other divisions and use the same terminology as in Unit III. Tables 9 through 11 of Unit IV.C. cover BPPD actions.

TABLE 9.—BIOPESTICIDES AND POLLUTION PREVENTION DIVISION—MICROBIAL AND BIOCHEMICAL PESTICIDES; NEW PRODUCTS AND AMENDMENTS

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
B580	86	New active ingredient; food use; establish tolerance[1]	18	18	44,100
B590	87	New active ingredient; food use; establish tolerance exemption[1]	16	16	27,563

TABLE 9.—BIOPESTICIDES AND POLLUTION PREVENTION DIVISION—MICROBIAL AND BIOCHEMICAL PESTICIDES; NEW PRODUCTS AND AMENDMENTS—Continued

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
B600	88	New active ingredient; non-food use[1]	12	12	16,538
B610	89	Food use; Experimental Use Permit application; establish temporary tolerance exemption	9	9	11,025
B620	90	Non-food use; Experimental Use Permit application	6	6	5,513
B621	91	Extend or amend Experimental Use Permit	6	6	4,410
B630	92	First food use; establish tolerance exemption	12	12	11,025
B631	93	Amend established tolerance exemption	9	9	11,025
B640	94	First food use; establish tolerance[1]	18	18	16,538
B641	95	Amend established tolerance (e.g., decrease or increase)	12	12	11,025
B650	96	New use; non-food	6	6	5,513
B660	97	New product; identical or substantially similar in composition and use to a registered product; no data review or only product chemistry data; cite-all data citation, or selective data citation where applicant owns all required data, or applicant submits specific authorization letter from data owner. Category also includes 100% re-package of registered end-use or manufacturing-use product that requires no data submission nor data matrix.	3	3	1,103
B670	98	New product; registered source of active ingredient; all Tier I data for product chemistry, toxicology, non-target organisms, and product performance must be addressed with product-specific data or with request for data waivers supported by scientific rationales	6	6	4,410
B671	99	New product; food use; unregistered source of active ingredient; requires amendment of established tolerance or tolerance exemption; all Tier I data requirements for product chemistry, toxicology, non-target organisms, and product performance must be addressed with product-specific data or with request for data waivers supported by scientific rationales	16	16	11,025
B672	100	New product; non-food use or food use having established tolerance or tolerance exemption; unregistered source of active ingredient; no data compensation issues; all Tier I data requirements for product chemistry, toxicology, non-target organisms, and product performance must be addressed with product-specific data or with request for data waivers supported by scientific rationales	12	12	7,875
B680	101	Label amendment requiring data submission[2]	4	4	4,410
B681	102	Label amendment; unregistered source of active ingredient; supporting data require scientific review	6	6	5,250
B682	103	Protocol review; applicant-initiated; excludes time for HSRB review (pre-application)	3	3	2,100

[1]All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.

[2]EPA-initiated amendments shall not be charged fees. Fast-track amendments handled by the Antimicrobials Division are to be completed within the FIFRA stated timelines listed in section 3(h) and are not subject to PRIA fees. Label amendments submitted by notification under PR Notices, such as PR Notice 95–2 and PR Notice 98–10, continue under PR Notice timelines and are not subject to PRIA fees.

TABLE 10.—BIOPESTICIDES AND POLLUTION PREVENTION DIVISION—STRAIGHT CHAIN LEPIDOPTERAN PHEROMONES (SCLPS)

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
B690	104	New active ingredient; food or non-food use[1]	6	6	2,205
B700	105	Experimental Use Permit application; new active ingredient or new use	6	6	1,103
B701	106	Extend or amend Experimental Use Permit	3	3	1,103
B710	107	New product; identical or substantially similar in composition and use to a registered product; no data review or only product chemistry data; cite-all data citation, or selective data citation where applicant owns all required data, or applicant submits specific authorization letter from data owner. Category also includes 100% re-package of registered end-use or manufacturing-use product that requires no data submission nor data matrix.	3	3	1,103
B720	108	New product; registered source of active ingredient; all Tier I data for product chemistry, toxicology, non-target organisms, and product performance must be addressed with product specific data or with request for data waivers supported by scientific rationales	4	4	1,103
B721	109	New product; unregistered source of active ingredient	6	6	2,310
B722	110	New use and/or amendment to tolerance or tolerance exemption	6	6	2,310
B730	111	Label amendment requiring data submission[2]	4	4	1,103

[1]All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.

[2]EPA-initiated amendments shall not be charged fees. Fast-track amendments handled by the Antimicrobials Division are to be completed within the FIFRA stated timelines listed in section 3(h) and are not subject to PRIA fees. Label amendments submitted by notification under PR Notices, such as PR Notice 95–2 and PR Notice 98–10, continue under PR Notice timelines and are not subject to PRIA fees.

TABLE 11.—BIOPESTICIDE AND POLLUTION PREVENTION DIVISION—PLANT INCORPORATED PROTECTANTS (PIPS)

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
B740	112	Experimental Use Permit application; registered active ingredient; non-food/feed or crop destruct basis; no SAP review required[1]	6	6	82,688
B750	113	Experimental Use Permit application; registered active ingredient; establish temporary tolerance or tolerance exemption; no SAP review required[1]	9	9	110,250
B760	114	Experimental Use Permit application; new active ingredient; non-food/feed or crop destruct basis; SAP review required; credit $78,750 toward new active ingredient application that follows	12	12	137,813
B761	115	Experimental Use Permit application; new active ingredient; non-food/feed or crop destruct; no SAP review required; credit $78,750 toward new active ingredient application that follows	7	7	82,688

TABLE 11.—BIOPESTICIDE AND POLLUTION PREVENTION DIVISION—PLANT INCORPORATED PROTECTANTS (PIPS)—Continued

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
B770	116	Experimental Use Permit application; new active ingredient; establish temporary tolerance or tolerance exemption; SAP review required; credit $105,000 toward new active ingredient application that follows	15	15	165,375
B771	117	Experimental Use Permit application; new active ingredient; establish temporary tolerance or tolerance exemption; no SAP review required; credit $105,000 toward new active ingredient application that follows	10	10	110,250
B772	118	Amend or extend Experimental Use Permit; minor changes to experimental design; established temporary tolerance or tolerance exemption is unaffected	3	3	11,025
B773	119	Amend or extend existing Experimental Use Permit; minor changes to experimental design; extend established temporary tolerance or tolerance exemption	5	5	27,563
B860	120	Amend Experimental Use Permit; first food use or major revision of experimental design	6	6	11,025
B780	121	New active ingredient; non-food/feed; no SAP review required[2]	12	12	137,813
B790	122	New active ingredient; Non-food/feed; SAP review required[2]	18	18	192,938
B800	123	New active ingredient; establish permanent tolerance or tolerance exemption based on temporary tolerance or tolerance exemption; no SAP review required[2]	12	12	220,500
B810	124	New active ingredient; establish permanent tolerance or tolerance exemption based on temporary tolerance or tolerance exemption; SAP review required[2]	18	18	275,625
B820	125	New active ingredient; establish tolerance or tolerance exemption; no SAP review required[2]	15	15	275,625
B840	126	New active ingredient; establish tolerance or tolerance exemption; SAP review required[2]	21	21	330,750
B830	127	New active ingredient; Experimental Use Permit application submitted simultaneously; establish tolerance or tolerance exemption; no SAP review required[2]	15	15	330,750
B850	128	New active ingredient; Experimental Use Permit requested simultaneously; establish tolerance or tolerance exemption; SAP review required[2]	21	21	385,875
B851	129	New active ingredient; different genetic event of a previously approved active ingredient; same crop; no tolerance action required; no SAP review required	9	9	110,250
B852	130	New active ingredient; different genetic event of a previously approved active ingredient; same crop; no tolerance action required; SAP review required	9	9	165,375
B870	131	New use[1]	9	9	33,075
B880	132	New product; no SAP review required[3]	9	9	27,563

TABLE 11.—BIOPESTICIDE AND POLLUTION PREVENTION DIVISION—PLANT INCORPORATED PROTECTANTS (PIPS)—Continued

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
B881	133	New product; SAP review required[3]	15	15	82,688
B890	134	Amendment; seed production to commercial registration; no SAP review required	9	9	55,125
B891	135	Amendment; seed production to commercial registration; SAP review required	15	15	110,250
B900	136	Amendment (except #B890); no SAP review required; (e.g., new IRM requirements that are applicant-initiated; or amending a conditional registration to extend the registration expiration date with additional data submitted) [4]	6	6	11,025
B901	137	Amendment (except #B890); SAP review required[4]	12	12	66,150
B902	138	PIP protocol review	3	3	5,513
B903	139	Inert ingredient tolerance exemption; e.g., a marker such as NPT II; reviewed in BPPD	6	6	55,125
B904	140	Import tolerance or tolerance exemption; processed commodities/food only	9	9	110,250

[1]Example: Transfer existing PIP trait by traditional breeding, such as from field corn to sweet corn.
[2]May be either a registration for seed increase or a full commercial registration. If a seed increase registration is granted first, full commercial registration is obtained using B890.
[3]Example: Stacking PIP traits within a crop using traditional breeding techniques.
[4]EPA-initiated amendments shall not be charged fees. Fast-track amendments handled by the Antimicrobials Division are to be completed within the FIFRA stated timelines listed in section 3(h) and are not subject to PRIA fees. Label amendments submitted by notification under PR Notices, such as PR Notice 95–2 and PR Notice 98–10, continue under PR Notice timelines and are not subject to PRIA fees.

V. How to Pay Fees

Applicants must submit fee payments at the time of application, and EPA will reject any application that does not contain evidence that the fee has been paid. EPA has developed a web site at *http://www.epa.gov/pesticides/fees/tool/index.htm* to help applicants identify the fee category and the fee. All fees should be rounded up to the whole dollar. Payments may be made by check, bank draft, or money order or online with a credit card or wire transfer.

A. Online

You may pay electronically through the government payment website *www.pay.gov*.

1. From the pay.gov home page, under ''Find Public Forms.''
2. Select ''search by Agency name.''
3. On the A-Z Index of Forms page, select ''E.''
4. Select ''Environmental Protection Agency.''
5. From the list of forms, select ''Prepayment of Pesticide Registration Improvement Act Fee.''
6. Complete the form entering the PRIA fee category and fee.
7. Keep a copy of the pay.gov acknowledgement of payment. A copy of the acknowledgement must be printed and attached to the front of the application to assure that EPA can match the application with the payment.

B. By Check or Money Order

All payments must be in United States currency by check, bank draft, or money order drawn to the order of the Environmental Protection Agency. On the check, the applicant must supply in the information line either the registration number of the product or the company number. A copy of the check must accompany the application to the Agency, specifically attached to the front of the application. The copy of the check ensures that payment has been made at the time of application and will enable the Agency to properly connect the payment with the application sent to the Agency.

If you send the Agency a check, it will be converted into an electronic funds transfer (EFT). This means the Agency will copy your check and use the account information on it to electronically debit your account for the amount of the check. The debit from your account will usually occur within 24 hours and will be shown on your regular account statement.

You will not receive your original check back. The Agency will destroy your original check but will keep the copy of it. If the EFT cannot be processed for technical reasons, you authorize the Agency to process the copy in place of your original check. If the EFT cannot be completed because of insufficient funds, the Agency may try to make the transfer up to two times.

All paper-based payments should be sent to the following address:

1. *By U.S. Postal Service*. U.S. Environmental Protection Agency, Washington Finance Center, FIFRA Service Fees, P.O. Box 979074, St. Louis, MO 63197–9000.

2. *By courier or personal delivery*. U.S. Bank, Government Lockbox 979074, 1005 Convention Plaza, SL–MO–C2–GL, St. Louis, MO 63197, (314) 418–4990.

VI. How to Submit Applications

Submissions to the Agency should be made at the address given in Unit VII. The applicant should attach documentation that the fee has been paid which may be pay.gov payment acknowledgement or a copy of the check. If the applicant is applying for a fee waiver, the applicant should provide sufficient documentation as described

in FIFRA section 33(b)(7) and *http://www.epa.gov/pesticides/fees/questions/waivers.htm*. The fee waiver request should be easy to identify and separate from the rest of the application and submitted with documentation that at least 25% of the fee has been paid.

If evidence of fee payment (electronic acknowledgement or copy of check properly identified as to company) is not submitted with the application, EPA will reject the application and will not process it further.

After EPA receives an application and payment, EPA performs a screen on the application to determine that the category is correct and that the proper fee amount has been paid. If either is incorrect, EPA will notify the applicant and require payment of any additional amount due. A refund will be provided in case of an overpayment. EPA will not process the application further until the proper fee has been paid for the category of application or a request for a fee waiver accompanies the application and the appropriate portion of the fee has been paid.

EPA will assign a unique identification number to each covered application for which payment has been made. EPA notifies the applicant of the unique identification number. This information is sent by e-mail if EPA has either an e-mail address on file or an e-mail address is provided on the application.

VII. Addresses

New covered applications should be identified in the title line with the mail code REGFEE.

1. *By USPS mail*. Document Processing Desk (REGFEE), Office of Pesticide Programs (7504P), U.S. Environmental Protection Agency, 1200 Pennsylvania Avenue, NW., Washington, DC 20460–0001.

2. *By courier*. Document Processing Desk (REGFEE), Office of Pesticide Programs, U.S. Environmental Protection Agency, Room S–4400, One Potomac Yard (South Bldg.), 2777 S. Crystal Drive, Arlington, VA 22202–4501.

Couriers and delivery personnel must present a valid picture identification card to gain access to the building. Hours of operation for the Document Processing Desk are 8 a.m. to 4:30 p.m., Monday through Friday, excluding Federal holidays.

List of Subjects

Environmental protection, Administrative practice and procedure, Pesticides.

Dated: July 28, 2008.

Marty Monell,

Acting Director, Office of Pesticide Programs.

[FR Doc. E8–17936 Filed 8–4–08; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPA–2008–0563 FRL–8701–1]

Relocation of National Oil and Hazardous Substance Pollution Contingency Plan, Subpart J Confidential Business Information Files

AGENCY: Environmental Protection Agency.

ACTION: Notice of Availability.

SUMMARY: The Environmental Protection Agency (EPA) Office of Solid Waste and Emergency Response, Office of Emergency Management announces that the Confidential Business Information (CBI) for products listed on the National Oil and Hazardous Substance Pollution Contingency Plan, under 40 CFR 300, Subpart J will be moved to a new contractor's office location. System Research and Applications Corporation (SRA) was awarded the new contract to support work on 40 CFR 300, Subpart J. The CBI files will be moved from the Computer Sciences Corporation (CSC) office in Alexandria, VA to SRA's office in Arlington, VA.

DATES: Comments must be received on or before August 11, 2008. The CBI files will be moved from the Computer Sciences Corporation (CSC) office in Alexandria, VA to SRA's office in Arlington, VA on Tuesday, August 12, 2008.

ADDRESSES: Submit your comments, identified by Docket ID No. EPA–HQ–OPA–2008–0563 by one of the following methods:

- *http://www.regulations.gov:* Follow the on-line instructions for submitting comment.
- *Mail:* The mailing address of the docket for this rule making is the EPA Docket Center (EPA/DC), Docket ID No. EPA–HQ–OPA–2008–0563, EPA West, Room 3334, 1301 Constitution Ave., NW., Washington, DC 20460.
- *Hand Delivery:* Such deliveries are only accepted during the Docket's normal hours of operation, and special arrangements should be made for deliveries of boxed information.

Instructions: Direct your comments to Docket ID No. EPA–HQ–OPA–2008–0563. EPA's policy is that all comments received will be included in the public docket without change and may be made available online at *http://www.regulations.gov,* including any personal information provided, unless the comment includes information claimed to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. Do not submit information that you consider to be CBI or otherwise protected through *http://www.regulations.gov* or *superfund.docket@epa.gov*. The *http://www.regulations.gov* Web site is an ''anonymous access'' system, which means EPA will not know your identity or contact information unless you provide it in the body of your comment. If you send an e-mail comment directly to EPA without going through *http://www.regulations.gov* your e-mail address will be automatically captured and included as part of the comment that is placed in the public docket and made available on the Internet. If you submit an electronic comment, EPA recommends that you include your name and other contact information in the body of your comment and with any disk or CD–ROM you submit. If EPA cannot read your comment due to technical difficulties and cannot contact you for clarification, EPA may not be able to consider your comment. Electronic files should avoid the use of special characters, any form of encryption, and be free of any defects or viruses. For additional information about EPA's public docket visit the EPA Docket Center homepage at *http://www.epa.gov/epahome/dockets.htm.*

Docket: All documents in the docket are listed in the *http://www.regulations.gov* index. Although listed in the index, some information is not publicly available, e.g., CBI or other information whose disclosure is restricted by statute. Certain other material, such as copyrighted material, will be publicly available only in hard copy. Publicly available docket materials are available either electronically in *http://www.regulations.gov* or in hard copy form at the EPA Docket Center, EPA/DC, EPA West, Room 3334, 1301 Constitution Ave., NW., Washington, DC. The Public Reading Room is open from 8:30 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. The telephone number for the Public Reading Room is (202) 566–1744, and the telephone number for the EPA Docket Center, EPA/DC is (202) 566–0276.

FOR FURTHER INFORMATION CONTACT: Leigh DeHaven, Office of Solid Waste and Emergency Response, Office of Emergency Management (5104A),